SP 02 200 3

FORM PTO - 1449

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

ATTORNEY DOCKET NO.: PXE-001C1

APPLICANT(S): Boyd et al.

SERIAL NO.: 10/764,328

FILING DATE: January 23, 2004 GROUP: 1614

					FILING	DATE: Ja	nuary 23,	2004 G	KOUI	P: 1614			
			U	.S. PATENT	DOCUM	ENTS							
EXAM. INIT.		DOCUMENT NUMBER	DATE	NAME	NAME		CLASS	SUB CLASS	FILING DATE IF APPROPRIATE				
			1										
			FOR	EIGN PATE	VT DOC	UMENT	3			· · · · · · · · · · · · · · · · · · ·			
EXAM. INIT.		DOCUMENT NUMBER	1 1 1		SUB CLASS	FILING DATE	ABSTRACT ENGLISH ONLY LANG (Y/N)						
				<u> </u>									
			OTHER	ART, JOURN	NAL AR	ricles,	ETC.	·					
EXAM. INIT.	ОТН	ER DOCUMEN	TS: (Includ	ling Author, Tit	tie, Date, F	lelevant Pa	ges, Place	of Publicat	ion)				
/CW/	C6	Bacchelli et al., (1999), "Identification of Heterozygote Carriers in Families with a Recessive Form of Pseudoxanthoma Elasticum (PXE)," Modern Pathology, 12(12):1112-1123.											
	C7	C7 Bergen et al., (2000), "Mutations in ABCC6 cause pseudoxanthoma elasticum," Nature Genetics, 25(2): 231.						<u>cs,</u> 25(2):228					
	C8 Cai et al., (2000), "A 500-kb region on chromosome 16p1 high-resolution mapping and genomic structure," <u>Journal</u>			-									
	C9 Germain et al., (2000), "Homozygosity for the R1268Q Mutation in MRP6, the Pseudoxanthoma Elast Gene, Is Not Disease-Causing," <u>Biochemical and Biophysical Research Communications</u> , 274(2):297- C10 Germain et al., (2000), "Identification of Two Polymorphisms (c189G>C; c190T>C) in Exon 2 of the Human MRP6 Gene (ABCC6) by Screening of Pseudoxanthoma Elasticum Patients: Possible Sequence Correction?" <u>Human Mutation</u> , Mutation and Polymorphism Report #174, Online (3 pages).												
	C11 Hopper et al., (2001), "Analysis of the structure and expression pattern of MRP7 (ABCC10), a new r of the MRP subfamily," Cancer Letters, 162(2):181-191.							new member					
	C12 Kool et al., (1999), "Expression of Human MRP6, a Homologue of the Multidrug Resistance Protei MRP1, in Tissues and Cancer Cells," Cancer Research, 59:175-182.							rotein Gene					
	C13 Le Saux et al., (2000), "Mutations in a gene encoding an ABC Transporter cause pseudoxanthoma elasticum," Nature Genetics, 25: 223-227.							oma					
	C14	Le Saux et al., (2001), "A Spectrum of ABCC6 Mutations Is Responsible for Pseudoxanthoma Elasticum," <u>American Journal of Human Genetics</u> , 69: 749-764.											
	C15	1		e activity of puri " <u>Biochimica et</u>			_	esistance p	rotein	MRP1 from			
EXAMINER		/Cy	ynthia Wil	der/	DATE	CONSIDE	RED	11/2	6/200)7			

FORM P	TO – 1	449	ATTORNEY DOCKET NO.: PXE-001C1						
		AL INFORMATION	APPLICANT(S): Boyd et al.						
DISCLO	SURE	STATEMENT	SERIAL NO.: 10/764,328						
			FILING DATE: January 23, 2004 GROUP: 1614						
OTHER ART, JOURNAL ARTICLES, ETC.									
EXAM. INIT.	·								
/CW/	C16 Perdu et al., (2000), "Identification of Novel Polymorphisms in the pM5 and MRP1 (ABCCI) Genes at Locus 16p13.1 and Exclusion of Both Genes as Responsible for Pseudoxanthoma Elasticum," Human Mutation, Mutation in Brief #388, Online (5 pages).								
	C17	Quaglino et al., (2000), "Abnormal phenotype of in vitro dermal fibroblasts from patients with pseudoxanthoma elasticum (PXE)," <u>Biochimica et Biophysica Acta</u> , 1501:51-62.							
	C18	Renie et al., (1984), "Pseudoxanthoma Elasticum: High Calcium Intake in Early Life Correlates With Severity," American Journal of Medical Genetics, 19:235-244.							
	C19	Ringpfeil et al., (2000), "Mutations in the MRP6 Gene Cause Pseudoxanthoma Elasticum," The Journal of Investigative Dermatology, 115(2):332, Abstract HB1.							
	C20	Sapadin et al., (1998), "Periumbilical pseudoxanthoma elasticum associated with chronic renal failure and angioid streaks – apparent regression with hemodialysis," <u>Journal of the American Academy of Dermatology</u> , 39(2):338-344.							
	C21	Struk et al., (2000), "Mutations of the gene encoding the transmembrane transporter protein ABC-C6 cause pseudoxanthoma elasticum," <u>Journal of Molecular Medicine</u> , 78(5):282-286.							
	C22	Tybulewicz et al., (1991), "Neonatal Lethality and Lymphopenia in Mice with a Homozygous Disruption of the c-abl Proto-Oncogene," Cell, 65:1153-1163.							
EXAMIN	ER	/Cynthia Wilder/	DATE CONSIDERED 11/26/2007						